## **Amendments to the Claims.**

This listing of claims will replace all prior versions, and listings, of claims in the application.

## **Listing of Claims:**

1. (Currently amended) A method of producing a radiolabelled gallium complex by reacting a Ga<sup>3+</sup> radioisotope in a suitable solvent with a macrocyclic bifunctional chelating agent, wherein said macrocyclic bifunctional chelating agent is linked to a targeting vector selected from the group consisting of proteins, glycoproteins, lipoproteins, polypeptides, glycopolypeptides, lipopolypeptides, peptides, glycopeptides, lipopolypeptides, carbohydrates, nucleic acids, oligonucleotides or a part, a fragment, a derivative or a complex of the aforesaid compounds and small organic molecules;

characterised in that the reaction is carried out using microwave activation at 80 to 120 W for 20 s to 2 min.

- 2. (Previously presented) The method according to claim 1 wherein the Ga<sup>3+</sup> radioisotope is selected from the group consisting of <sup>66</sup>Ga<sup>3+</sup>, <sup>67</sup>Ga<sup>3+</sup> and <sup>68</sup>Ga<sup>3+</sup>.
- 3. (Previously presented) The method according to claim 1 wherein the Ga<sup>3+</sup> radioisotope is <sup>68</sup>Ga<sup>3+</sup>.
- 4. (Cancelled)
- 5. (Currently amended) The method according to claim 1 wherein the <u>macrocyclic</u> <u>bifunctional</u> chelating agent comprises hard donor atoms, preferably O and N atoms.
- 6. (Cancelled)

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7. (Cancelled)

8. (Currently amended) The method according to claim 17 wherein the target vector is a

peptide or oligonucleotide.

9. (Previously presented) The method according to claim 1 wherein the microwave

activation is carried out at 90 to 110 W.

10. (Previously presented) The method according to claim 1 wherein the microwave

activation is carried out for 30 s to 90 s.

11. (Previously presented) The method according to claim 3 wherein the <sup>68</sup>Ga<sup>3+</sup> is obtained

by contacting the eluate from a <sup>68</sup>Ge/<sup>68</sup>Ga generator with an anion exchanger and eluting

<sup>68</sup>Ga<sup>3+</sup> from said anion exchanger.

12. (Previously presented) The method according to claim 11 wherein the <sup>68</sup>Ge/<sup>68</sup>Ga

generator comprises a column comprising titanium dioxide.

13. (Previously presented) The method according to claim 11 wherein the anion exchanger

comprises HCO<sub>3</sub> as counterions.

14. (Previously presented) The method according to claim 11 wherein the anion exchanger is

an anion exchanger comprising quaternary amine functional groups, or the ion exchanger

is a anion exchange resin based on polystyrene-divinylbenzene.

15. (Previously presented) The method according to claim 1 for the production of <sup>68</sup>Ga-

radiolabelled PET tracers.

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16. (Withdrawn) Method according to claim 11 wherein the eluting <sup>68</sup>Ga<sup>3+</sup> is in the picomolar to nanomolar range after the elution, and more preferably in a nanomolar to micromolar level.